

JAVALI A

Hyderabad, India | Phone: +91 9441610954 | Mail: javali.adiraju@gmail.com | www.linkedin.com/in/javali-adiraju

CAREER OBJECTIVE

- Pharm.D graduate with strong foundation in ICH-GCP, pharmacology, clinical medicine, and therapeutics. Basics in Good Pharmacovigilance Practices (GVP), Good Clinical Practice (GCP). Seeking a challenging entry-level role to apply scientific acumen and communication skills in cross-functional environments.

EDUCATIONAL QUALIFICATION

- Doctor Of Pharmacy** Hyderabad, India
Sept 2019- Aug 2025
MNR hospital affiliated to **Osmania university** , Percentage: 78%
Hands-on exposure to: ADR monitoring & reporting, Antibiotic audit, Drug interaction assessment & report, Prescription audit & medication error monitoring.
- Sri Gayathri Junior College** Hyderabad, India
July 2015 - May2017
Senior Secondary Education
- Triveni Talent School** Hyderabad, India
May 2015
Senior Secondary Education

EXPERIENCE

Pharm D Intern

KIMS-Sunshine Hospitals, Begumpet – Hyderabad Duration: 3 Months

- Gained practical experience in clinical documentation, Document-level Quality Control (QC). Maintained Patient Safety & Data Integrity. Participated in patient counseling and medication reconciliation rounds.
- Performed Patient safety monitoring by performing life case audits, and monitored the therapies for errors in drug doses, ADRs interactions in prescription audits. Coordinated with cross functional teams like nurses, physiotherapists, dietitians in the view of patient safety and counseling.

ACADEMIC PROJECT

Title: Assessment Of Maternal And Neonatal Outcomes Among Women At High Risk Preeclampsia And Normotensive Pregnancy.

- Developed clinical study protocol and following ICH- GCP guidelines under guidance.
- Performed **literature searches** for safety sections using **PubMed and Google Scholar** and other search engines. Gained exposure to clinical trial documentation practices compliant with ICH-GCP guidelines.
- Performed quality checks and assist in study document review and expressed the clinical data in TFLs (Tables, Figures, Listings).
- Management of references and citation in Vancouver style is done by using reference software tool **ZOTERO & EndNote**, Thesis written in compliance with industry standards.

CERTIFICATIONS

- **Certified in Good Clinical Practice (GCP)** – National Institute on Drug Abuse (NIDA).
- **Certified in Basic course in biomedical research** – Indian council of medical research (ICMR), National institute of epidemiology (NIE).
- **Certified in Data analysis** – Swayam.
- **Certified in Master Pharma Regulatory Affairs: FDA/EMA & Global** – Udemy.
- **Certified in Medical Writing**- Unisearch Life sciences.
- **Certified in Pharmacovigilance Basics-1** – VIGISERVE Foundation.
- **Oral Presenter** – JNTU University College of Pharmaceutical Sciences, Sulthanpur.

PUBLICATIONS

- Assessment Of Maternal and Neonatal Outcomes Among Women at High-risk Pre-eclampsia and Normotensive Pregnancy. (2024). International Journal of Medical Science and Clinical Invention, 11(07), 7177-7184. <https://doi.org/10.18535/ijmsci/v11i7.01>.**

TECHNICAL SKILLS

- Medical writing.
- CTD 5 modules.
- Clinical trial phases, documentation in clinical trial.
- Basics in drug development process, ICH-GCP, and regulatory guidelines.
- Knowledge on medical terminology and therapeutic areas.
- Strong knowledge in Microsoft Word, Excel & PowerPoint.

SOFT SKILLS

- Attention to detail.
- Critical thinking & problem-solving skills.
- Strong written and verbal English communication skills.
- Adaptability & Flexibility.
- Effective communication with cross-functional teams.
- Team player, Time management and deadline-oriented.

DECLARATION

I hereby declare that all the particulars mentioned above are true to the best of my knowledge and belief.

Place: Sangareddy

Javali A